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## **ABSTRACT**

Angiogenesis inhibiting 5-substituted-1,2,4-thiadiazolyl derivatives

This invention concerns compounds of formula

$$\begin{array}{c|c}
 & R^2 \\
 & R^3 \\
 & R^4
\end{array}$$
(I),

the N-oxide forms, the pharmaceutically acceptable acid addition salts and stereochemically isomeric forms thereof, wherein X is CH or N; R<sup>1</sup> is hydrogen,  $C_{1-6}$ alkyl,  $C_{1-6}$ alkyloxy,  $C_{1-6}$ alkylthio, amino, mono- or di( $C_{1-6}$ alkyl)amino,  $Ar^{1}$ , Ar<sup>1</sup>NH-, C<sub>3-6</sub>cycloalkyl, hydroxymethyl or benzyloxymethyl; R<sup>2</sup> is hydrogen, 15 C<sub>1-6</sub>alkyl, amino, aminocarbonyl, mono- or di(C<sub>1-6</sub>alkyl)amino, C<sub>1-6</sub>alkyloxycarbonyl, C<sub>1-6</sub>alkylcarbonylamino, hydroxy or C<sub>1-6</sub>alkyloxy; R<sup>3</sup>, R<sup>4</sup> and R<sup>5</sup> are each independently selected from hydrogen, halo, C<sub>1-6</sub>alkyl, C<sub>1-6</sub>alkyloxy, trifluoromethyl, nitro, amino, cyano, azido, C<sub>1-6</sub>alkyloxyC<sub>1-6</sub>alkyl, C<sub>1-6</sub>alkylthio, C<sub>1-</sub> 6alkyloxycarbonyl or  $Het^1$ ; —(A)— is  $Ar^2$ ,  $Ar^2CH_2$ - or  $Het^2$ ;  $Ar^1$  and  $Ar^2$  optionally 20 substituted phenyl; Het1 and Het2 are optionally substituted monocyclic heterocycles; having angiogenesis inhibiting activity; their preparation, compositions containing

them and their use as a medicine.